

=> fil reg; d stat que 18; fil capl uspatf toxcenter; s 18
FILE 'REGISTRY' ENTERED AT 10:17:03 ON 31 MAR 2006
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Property values tagged with IC are from the ZIC/VINITI data file provided by InfoChem.

STRUCTURE FILE UPDATES: 29 MAR 2006 HIGHEST RN 878540-28-4
DICTIONARY FILE UPDATES: 29 MAR 2006 HIGHEST RN 878540-28-4

New CAS Information Use Policies, enter HELP USAGETERMS for details.

TSCA INFORMATION NOW CURRENT THROUGH January 6, 2006

Please note that search-term pricing does apply when conducting SmartSELECT searches.

*

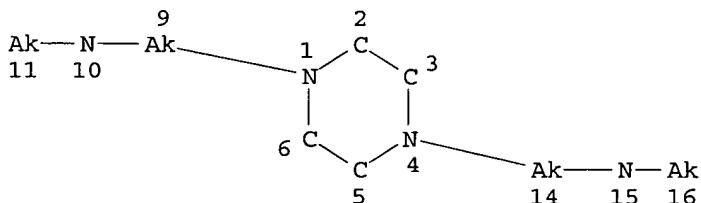
* The CA roles and document type information have been removed from *
* the IDE default display format and the ED field has been added, *
* effective March 20, 2005. A new display format, IDERL, is now *
* available and contains the CA role and document type information. *

Structure search iteration limits have been increased. See HELP SLIMITS for details.

REGISTRY includes numerically searchable data for experimental and predicted properties as well as tags indicating availability of experimental property data in the original document. For information on property searching in REGISTRY, refer to:

<http://www.cas.org/ONLINE/UG/regprops.html>

L6 STR



Ak = alkyl

NODE ATTRIBUTES:

CONNECT IS E2 RC AT 9

CONNECT IS E1 RC AT 11

CONNECT IS E2 RC AT 14

CONNECT IS E1 RC AT 16

DEFAULT MLEVEL IS ATOM

MLEVEL IS CLASS AT 9 11 14 16

GGCAT IS LIN LOC SAT AT 9

GGCAT IS UNS AT 11

GGCAT IS LIN LOC SAT AT 14

GGCAT IS UNS AT 16

 } alkyls at nodes 9 & 14 are linear, saturated,
 } with ≤ 6 carbons
 } alkyls at nodes 11 & 16 are unsaturated, with
 } ≥ 6 carbons

DEFAULT ECLEVEL IS LIMITED
ECOUNT IS M6 C AT 11
ECOUNT IS M6 C AT 16

GRAPH ATTRIBUTES:
RING(S) ARE ISOLATED OR EMBEDDED
NUMBER OF NODES IS 12

STEREO ATTRIBUTES: NONE
L8 1 SEA FILE=REGISTRY SSS FUL L6

100.0% PROCESSED 706610 ITERATIONS 1 ANSWERS
SEARCH TIME: 00.00.09

FILE 'CAPLUS' ENTERED AT 10:17:03 ON 31 MAR 2006
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COPYRIGHT (C) 2006 AMERICAN CHEMICAL SOCIETY (ACS)

FILE 'USPATFULL' ENTERED AT 10:17:03 ON 31 MAR 2006
CA INDEXING COPYRIGHT (C) 2006 AMERICAN CHEMICAL SOCIETY (ACS)

FILE 'TOXCENTER' ENTERED AT 10:17:03 ON 31 MAR 2006
COPYRIGHT (C) 2006 ACS

L13 4 L8

=> fil marpat; d stat que 112
FILE "MARPAT" ENTERED AT 10:17:20 ON 31 MAR 2006
USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT.
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FILE CONTENT: 1961-PRESENT VOL 144 ISS 10 (20060324/ED)

SOME MARPAT RECORDS ARE DERIVED FROM INPI DATA FOR 1961-1987

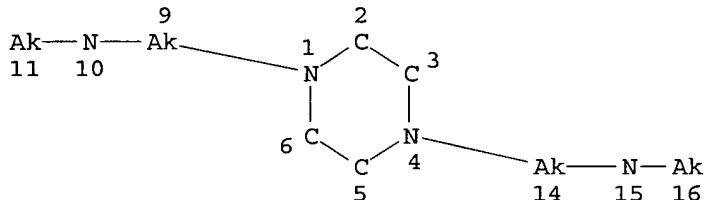
MOST RECENT CITATIONS FOR PATENTS FROM MAJOR ISSUING AGENCIES
(COVERAGE TO THESE DATES IS NOT COMPLETE):

US	2006035965	16 FEB 2006
DE	102004031947	19 JAN 2006
EP	1614691	11 JAN 2006
JP	2006016369	19 JAN 2006
WO	2006012333	02 FEB 2006
GB	2416167	18 JAN 2006
FR	2873371	27 JAN 2006
RU	2267521	10 JAN 2006
CA	2472818	30 DEC 2005

Expanded G-group definition display now available.

New CAS Information Use Policies, enter HELP USAGETERMS for details.

L6 STR



NODE ATTRIBUTES:

CONNECT IS E2 RC AT 9
CONNECT IS E1 RC AT 11
CONNECT IS E2 RC AT 14
CONNECT IS E1 RC AT 16
DEFAULT MLEVEL IS ATOM
MLEVEL IS CLASS AT 9 11 14 16
GGCAT IS LIN LOC SAT AT 9
GGCAT IS UNS AT 11
GGCAT IS LIN LOC SAT AT 14
GGCAT IS UNS AT 16
DEFAULT ECLEVEL IS LIMITED
ECOUNT IS M6 C AT 11
ECOUNT IS M6 C AT 16

*Same structure
as was searched in Registry*

GRAPH ATTRIBUTES:

RING(S) ARE ISOLATED OR EMBEDDED
NUMBER OF NODES IS 12

STEREO ATTRIBUTES: NONE

L12 15 SEA FILE=MARPAT SSS FUL L6

100.0% PROCESSED 29452 ITERATIONS
SEARCH TIME: 00.00.25

15 ANSWERS

=> dup rem l13,l12
FILE 'CAPLUS' ENTERED AT 10:17:30 ON 31 MAR 2006
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FILE 'USPATFULL' ENTERED AT 10:17:30 ON 31 MAR 2006
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FILE 'MARPAT' ENTERED AT 10:17:30 ON 31 MAR 2006
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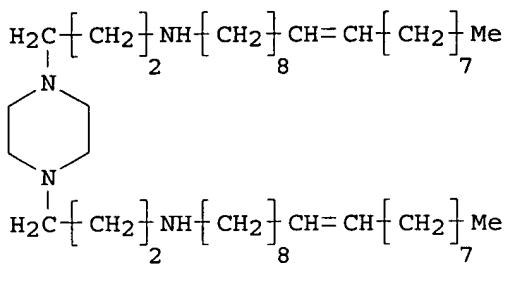
PROCESSING COMPLETED FOR L13
PROCESSING COMPLETED FOR L12

L14 16 DUP REM L13 L12 (3 DUPLICATES REMOVED)
ANSWER '1' FROM FILE CAPLUS
ANSWER '2' FROM FILE USPATFULL
ANSWERS '3-16' FROM FILE MARPAT

=> d ibib ed abs hitstr 1-2; d ibib ed abs qhit 3-16; fil hom

L14 ANSWER 1 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN DUPLICATE 1
ACCESSION NUMBER: 1998:263206 CAPLUS
DOCUMENT NUMBER: 128:266964
TITLE: Process of transfecting a cell with a polynucleotide
mixed with an amphipathic compound and a DNA-binding
protein
INVENTOR(S): Wolff, Jon A.; Fritz, Jeffery; Budker, Vladimir;
Hagstrom, James
PATENT ASSIGNEE(S): Mirus Corporation, USA
SOURCE: U.S., 16 pp.
CODEN: USXXAM
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 5744335	A	19980428	US 1995-530598	19950919
US 6180784	B1	20010130	US 1998-20566	19980117
PRIORITY APPLN. INFO.:			US 1995-530598	A3 19950919
OTHER SOURCE(S):	MARPAT	128:266964		
ED Entered STN:	08 May 1998			
GI				



AB Transfection of a cell is accomplished using with a polynucleotide mixed with one or more amphipathic compds. and a DNA-binding protein, especially a histone such as histones H1, H2A, or H2B. The DNA-binding protein may be fused to a nuclear localization signal peptide. Exemplary and preferred amphipathic compds. are cationic amphipathic compds. I was synthesized in 70% yield by reacting 1,4-bis(3-aminopropyl)piperazine with oleoyl chloride and reducing the intermediate with LiAlH4 in THF. Histone H1 was found to increase the transfection efficiency of I 16.1-fold. I/H1 reagent has a greater transfection efficiency and less cellular toxicity than LipofectAmine, which is useful in gene therapy.

IT 205595-99-9P

RL: BUU (Biological use, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

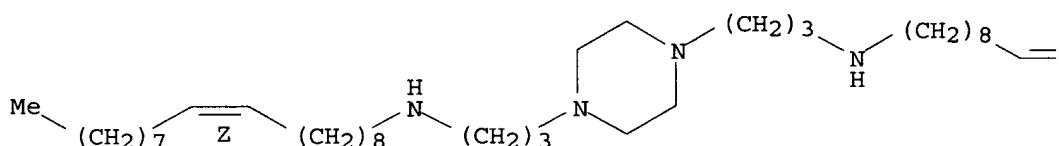
(cell transfection with polynucleotide mixed with amphipathic compound and DNA-binding protein)

RN 205595-99-9 CAPLUS

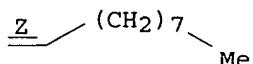
CN 1,4-Piperazinedipropanamine, N,N'-di-9-octadecenyl-, (Z,Z)- (9CI) (CA INDEX NAME)

Double bond geometry as shown.

PAGE 1-A



PAGE 1-B



REFERENCE COUNT: 25 THERE ARE 25 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L14 ANSWER 2 OF 16 USPATFULL on STN

ACCESSION NUMBER: 2001:14639 USPATFULL

TITLE: Process of transfecting a cell with a polynucleotide mixed with an amphipathic compound and a DNA-binding protein

INVENTOR(S): Wolff, Jon A., Madison, WI, United States

PATENT ASSIGNEE(S) : Hagstrom, James E., Madison, WI, United States
 Budker, Vladimir G., Madison, WI, United States
 Fritz, Jeffery, Nashville, TN, United States
 Mirus Corporation, Madison, WI, United States (U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 6180784	B1	20010130
APPLICATION INFO.:	US 1998-20566		19980117 (9)
RELATED APPLN. INFO.:	Division of Ser. No. US 1995-530598, filed on 19 Sep 1995, now patented, Pat. No. US 5744335		
DOCUMENT TYPE:	Utility		
FILE SEGMENT:	Granted		
PRIMARY EXAMINER:	Lambkin, Deborah C.		
LEGAL REPRESENTATIVE:	Johnson, Mark K.		
NUMBER OF CLAIMS:	8		
EXEMPLARY CLAIM:	1		
LINE COUNT:	1198		

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The present invention provides a process of transfecting a cell with a polynucleotide mixed with one or more amphipathic compounds and an effective amount of a DNA-binding protein. Exemplary and preferred DNA-binding proteins are H1, H2A, and H2B. Exemplary and preferred amphipathic compounds are cationic amphipathic compounds.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

IT 205595-99-9P

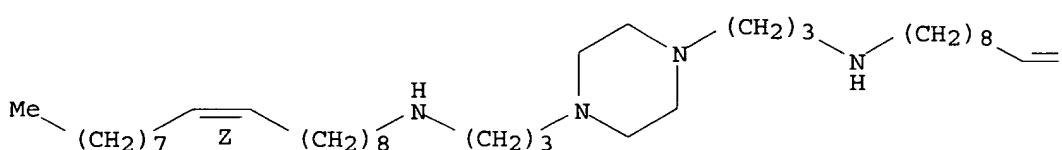
(cell transfection with polynucleotide mixed with amphipathic compound and DNA-binding protein)

RN 205595-99-9 USPATFULL

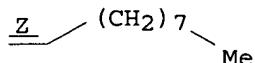
CN 1,4-Piperazinedipropanamine, N,N'-di-9-octadecenyl-, (Z,Z)- (9CI) (CA INDEX NAME)

Double bond geometry as shown.

PAGE 1-A



PAGE 1-B



'ED' IS NOT A VALID FORMAT

REENTER DISPLAY FORMAT FOR ALL FILES (FILEDEFAULT):ibib abs qhit

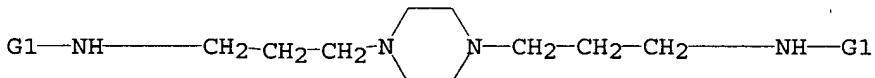
L14 ANSWER 3 OF 16 MARPAT COPYRIGHT 2006 ACS on STN
 ACCESSION NUMBER: 143:103230 MARPAT

TITLE: Compositions and processes using siRNA, amphipathic compounds, and polycations
 INVENTOR(S): Monahan, Sean D.; Lewis, David L.; Herweijer, Hans; Wolff, Jon A.; Hagstrom, James E.; Loomis, Aaron G.; Trubetskoy, Vladimir; Higgs, Lori
 PATENT ASSIGNEE(S): USA
 SOURCE: U.S. Pat. Appl. Publ., 34 pp., Cont.-in-part of U.S. Ser. No. 345,021.
 CODEN: USXXCO
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 33
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 2005143332	A1	20050630	US 2004-845968	20040514
US 2003125281	A1	20030703	US 2002-157674	20020528
US 2003143204	A1	20030731	US 2002-186757	20020701
US 2004137064	A1	20040715	US 2003-345021	20030115
WO 2005116045	A1	20051208	WO 2004-US15507	20040515
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
PRIORITY APPLN. INFO.:			US 2002-157674	20020528
			US 2002-186757	20020701
			US 2003-345021	20030115
			US 2001-917154	20010727
			US 2001-315394P	20010827
			US 2001-324155P	20010920
			US 2004-845968	20040514

AB Described is a deliverable composition with low toxicity comprising an amphipathic compound, a polycation, and a siRNA. The composition may be used in the process of delivering a siRNA to an animal cell or more particularly, a mammal cell.

MSTR 1

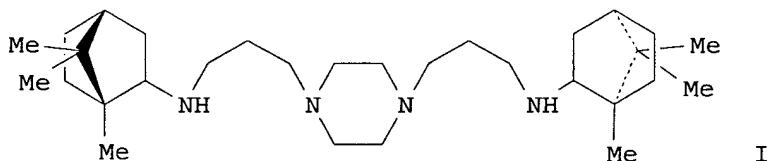


G1 = alkenyl <containing 6-24 C>
 Patent location: claim 3

L14 ANSWER 4 OF 16 MARPAT COPYRIGHT 2006 ACS on STN
 ACCESSION NUMBER: 143:440444 MARPAT
 TITLE: Preparation of polyamines as ligands for metal

complexes
 INVENTOR(S): Habagami, Shigeki; Higashimura, Hideyuki
 PATENT ASSIGNEE(S): Sumitomo Chemical Co., Ltd., Japan
 SOURCE: Jpn. Kokai Tokkyo Koho, 8 pp.
 CODEN: JKXXAF
 DOCUMENT TYPE: Patent
 LANGUAGE: Japanese
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 2005314273	A2	20051110	JP 2004-133185	20040428
PRIORITY APPLN. INFO.:			JP 2004-133185	20040428
GI				

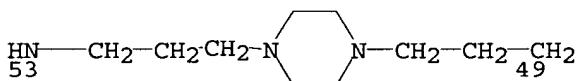


AB R1(NR1R2)nNR12 (n = 3-7; R1 = H, hydrocarbyl; R2 = hydrocarbylene; ≥2 of R1 hydrocarbyl having asym. C atom; 2 R2s may be bonded to form a ring; number of rings ≤ 3) are prepared. Thus, a mixture of (+)-camphor, toluene, BF₃-Et₂O, and 1,4-bis(3-aminopropyl)piperazine was refluxed for 18 h and the reaction product was treated with NiCl₂ and NaBH₄ at room temperature for 16 h to give 18% tetramine I. Oxidative polymerization of 2,3-dihydroxynaphthalene using I and CuCl gave 11% poly(2,3-dihydroxy-1,4-naphthalene).

MSTR 1

G1—G2—G7
 1 3

G1 = alkenyl <containing 2-30 C>
 G2 = 53-1 49-3



G5 = alkenyl <containing 2-30 C>
 G7 = 8

HN—G5
 8

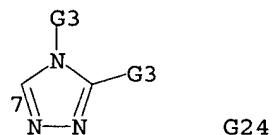
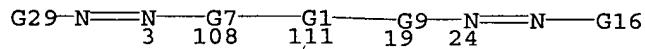
Patent location: claim 1
 Note: substitution is restricted
 Note: additional ring formation also claimed

L14 ANSWER 5 OF 16 MARPAT COPYRIGHT 2006 ACS on STN
 ACCESSION NUMBER: 144:93797 MARPAT
 TITLE: Particular diazo cationic compound, as direct dyes,
 for dyeing of hair
 INVENTOR(S): Greaves, Andrew; David, Herve
 PATENT ASSIGNEE(S): L'Oreal, Fr.
 SOURCE: Fr. Demande, 53 pp.
 CODEN: FRXXBL
 DOCUMENT TYPE: Patent
 LANGUAGE: French
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
FR 2872162	A1	20051230	FR 2004-6871	20040623
JP 2006009017	A2	20060112	JP 2005-182483	20050622
CN 1737064	A	20060222	CN 2005-10091353	20050622
US 2006021162	A1	20060202	US 2005-159154	20050623
EP 1634926	A1	20060315	EP 2005-291353	20050623
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, PL, SK, BA, HR, IS, YU				
PRIORITY APPLN. INFO.:			FR 2004-6871	20040623
			US 2004-588041P	20040715

AB Diazo cationic compds. are used as direct dyes for dyeing of hair (Markush structure given). A diazo cationic dye was prepared by the reaction of a diazonium aminopyridine derivative with an azopyridinium derivative. A hair dye containing 5x10⁻³ mol/L of above dye was applied to a 90% gray hair to obtain a fuchsia color.

MSTR 1



G1 = 131-108 133-19 / 145-108 149-19

G21—G22—G21 G21—G32—G21—G33—G21
 131 133 145 147 149

G7 = 17-3 18-111

G28—G8
 17 18

G8 = 28

N---G13
28

G9 = 26-111 27-24

G12-G14
26 27

G12 = 129

N---G13
129

G13 = carbon chain <containing 1-20 C> (opt. substd.)
 G21 = carbon chain <containing 1-14 C> (opt. substd.)
 G22 = 142-131 139-133



G31 = (1-2) CH₂

Patent location:

claim 1

Note: additional ring formation also claimed

Note: substitution is restricted

REFERENCE COUNT: 3 THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L14 ANSWER 6 OF 16 MARPAT COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 141:111545 MARPAT

TITLE: Compositions and processes using siRNA, amphipathic compounds, and polycations

INVENTOR(S): Lewis, David L.; Herweijer, Hans; Monahan, Sean D.; Wolff, Jon A.; Hagstrom, James E.; Loomis, Aaron G.

PATENT ASSIGNEE(S): USA

SOURCE: U.S. Pat. Appl. Publ., 23 pp.

CODEN: USXXCO

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 33

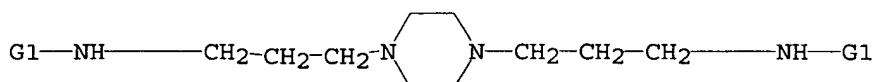
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 2004137064	A1	20040715	US 2003-345021	20030115
WO 2004065587	A1	20040805	WO 2003-US2165	20030124
W: JP				
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, SE, SI, SK, TR				
EP 1597357	A1	20051123	EP 2003-707518	20030124
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, FI, CY, TR, BG, CZ, EE, HU, SK				
US 2004019008	A1	20040129	US 2003-621760	20030717
US 2005143332	A1	20050630	US 2004-845968	20040514
PRIORITY APPLN. INFO.:			US 2002-157674	20020528

US 2002-186757 20020701
 US 2003-345021 20030115
 WO 2003-US2165 20030124

AB Described is a composition with low toxicity comprising an amphipathic compound and a polycation, useful for delivering siRNA to a cell. The composition may be used in the process of delivering a siRNA to an animal cell, or more particularly a mammal cell, in a multi-well format.

MSTR 1



G1 = alkenyl <containing 6-24 C>

Patent location: claim 4

L14 ANSWER 7 OF 16 MARPAT COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 140:184420 MARPAT

TITLE: Cyclic aminothioureas as lubricating oil antiwear, anticorrosion, and antioxidant additives to replace zinc dialkyl dithiophosphates

INVENTOR(S): Mukkamala, Ravindranath

PATENT ASSIGNEE(S): USA

SOURCE: U.S. Pat. Appl. Publ., 5 pp.

CODEN: USXXCO

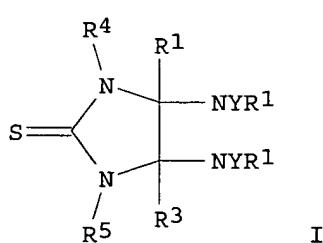
DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

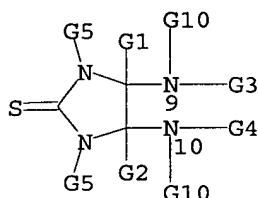
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 2004029746	A1	20040212	US 2003-636683	20030807
EP 1394243	A1	20040303	EP 2003-254693	20030728
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
JP 2004131710	A2	20040430	JP 2003-287847	20030806
PRIORITY APPLN. INFO.:			US 2002-401845P	20020807
GI				

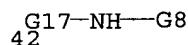


AB Cyclic and bicyclic aminothiourea-based lubricating oil antiwear, corrosion inhibitor, and antioxidant additives, are suitable replacements for zinc dialkyl dithiophosphates. The aminothioureas are of general structure I, in which R2 and R3 are H, alkyl, alkenyl, aryl, or aralkyl;

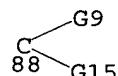
or R2R3 can combine with carbon atoms of an imidazolidinethione ring to form a saturated or unsatd. C5-8-carbocyclic ring; R1 is alkyl, alkenyl, or aralkyl; or R1 groups can combine with the nitrogen atoms to which they are attached and carbon atoms of an imidazolidinethione ring to form a five- to seven-membered heterocyclic ring. Further, R4 and R5 are H, alkyl, alkenyl, aryl aralkyl, -CHR6-CHR7-CO2R8, -CR9R10-NHR11, or -C(:Z)-NHR12; R6 and R7 are H or C1-4-alkyl; R8-10 are alkyl, alkenyl, aralkyl, or aryl; Y = H, alkyl, alkenyl, aralkyl, -CHR6-CHR7-CO2R8, -CR9R10-NHR11, -C(:Z)NHR12, or -OH; and Z = O or S. The additives, which can consist of 2 cyclic thioamides, are present at 0.1-20 weight% concentration

MSTR 1

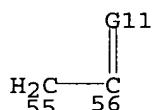
G8 = carbon chain <containing 1-22 C,
0 or more double bonds, no triple bonds> (opt. substd.)
G10 = 42



G17 = 88



G3 +G4 = 55-9 56-10



Patent location:
Note:

claim 1
oxygen alternative in G10 is free radical

L14 ANSWER 8 OF 16 MARPAT COPYRIGHT 2006 ACS on STN
ACCESSION NUMBER: 140:133819 MARPAT
TITLE: Compositions and processes using siRNA, amphipathic compounds and polycations
INVENTOR(S): Lewis, David L.; Hagstrom, James E.; Herweijer, Hans; Loomis, Aaron G.; Monahan, Sean D.; Wolff, Jon A.; Trubetskoy, Vladimir
PATENT ASSIGNEE(S): USA
SOURCE: U.S. Pat. Appl. Publ., 25 pp., Cont.-in-part of U.S. Ser. No. 345,021.

CODEN: USXXCO

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT:

33

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 2004019008	A1	20040129	US 2003-621760	20030717
US 2003125281	A1	20030703	US 2002-157674	20020528
US 2003143204	A1	20030731	US 2002-186757	20020701
US 2004137064	A1	20040715	US 2003-345021	20030115
WO 2005017098	A2	20050224	WO 2003-US25121	20030811
WO 2005017098	A3	20050630		

W: JP

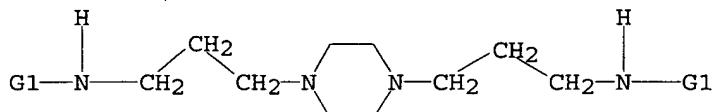
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE,
IT, LU, MC, NL, PT, RO, SE, SI, SK, TR

PRIORITY APPLN. INFO.:

US 2002-157674	20020528
US 2002-186757	20020701
US 2003-345021	20030115
US 2001-917154	20010727
US 2001-315394P	20010827
US 2001-324155P	20010920
US 2003-621760	20030717

AB Described is a composition with low toxicity comprising an amphipathic compound and a polycation, useful for delivering siRNA to a cell. The composition may be used in the process of delivering a siRNA to an animal cell, or more particularly a mammal cell, in a multi-well format.

MSTR 1



G1 = alkenyl <containing 6-24 C>
Patent location: claim 2

L14 ANSWER 9 OF 16 MARPAT COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 138:189529 MARPAT

TITLE: Ink sets with improved light discoloration resistance,
and method and apparatus for ink-jet recording
INVENTOR(S): Yakushigawa, Yuko; Teraoka, Hisashi; Mafune, Kumiko;
Kanke, Takeshi; Takizawa, Yoshihisa

PATENT ASSIGNEE(S): Canon Inc., Japan

SOURCE: Jpn. Kokai Tokkyo Koho, 35 pp.

CODEN: JKXXAF

DOCUMENT TYPE: Patent

LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 2

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 2003055585	A2	20030226	JP 2001-191882	20010625
US 6866380	B2	20050315	US 2003-682521	20031010

PRIORITY APPLN. INFO.:

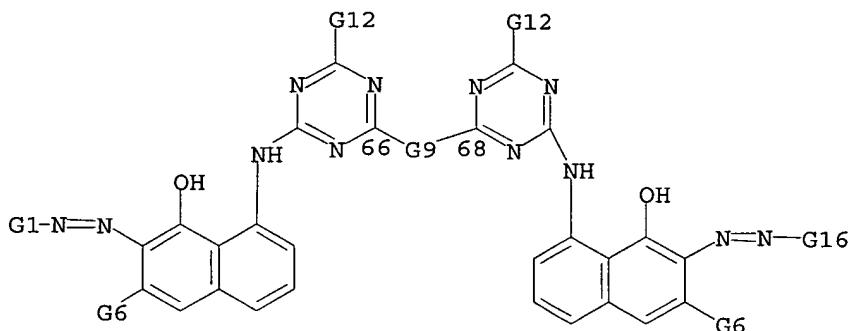
JP 2000-190325	20000623
JP 2001-168257	20010604
US 2001-884096	20010620

GI

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT *

AB The ink sets consist of ≥ 3 water-based color inks showing the difference of light discoloration ΔE (definition given) among the inks ≤ 10 and reflection d. retention after a fading test (condition given) $\geq 70\%$. Thus, an ink set was manufactured from a cyan ink containing C.I. Acid Blue 9 and C.I. Direct Blue 199, a magenta ink containing I, II, and C.I. Acid Red 289, and a yellow ink containing C.I. Direct Yellow 132.

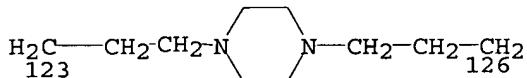
MSTR 3



G9 = 4-66 61-68

G14-G11-G14
4 61

G10 = alkenyl (opt. subst. by 1 or more G13)
 G11 = 123-4 126-61



G13 = CO₂H
 G14 = 74

N-G10
74

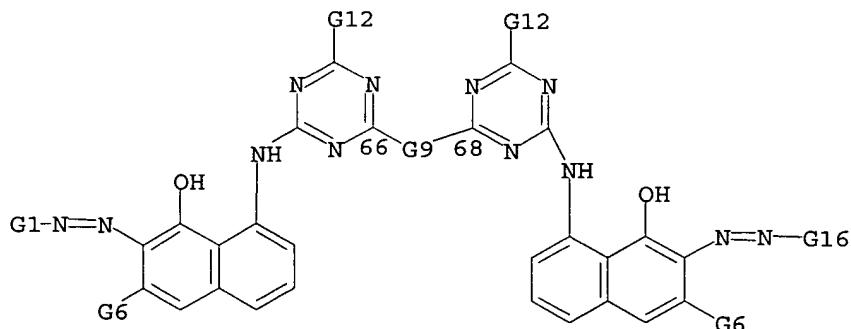
Patent location: claim 10
 Note: or salts
 Note: additional ring formation also claimed

L14 ANSWER 10 OF 16 MARPAT COPYRIGHT 2006 ACS on STN
 ACCESSION NUMBER: 138:138941 MARPAT
 TITLE: Light-resistant ink sets and apparatus and method for jet-printing
 INVENTOR(S): Mafune, Kumiko; Kanke, Takeshi
 PATENT ASSIGNEE(S): Canon Inc., Japan
 SOURCE: Jpn. Kokai Tokkyo Koho, 39 pp.
 CODEN: JKXXAF
 DOCUMENT TYPE: Patent
 LANGUAGE: Japanese
 FAMILY ACC. NUM. COUNT: 2
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 2003034763	A2	20030207	JP 2001-173046	20010607
US 2004074418	A1	20040422	US 2003-682519	20031010
PRIORITY APPLN. INFO.:			JP 2000-176138	20000612
			JP 2001-145161	20010515
			US 2001-871627	20010604

AB The ink sets comprise same color tone two water-thinned inks containing ≥ 1 common colorant, one of which shows lower color d. and gives images with the same or better light fastness. Thus, an ink containing 0.5 part C.I. Acid Blue 9 and 3.5 part C.I. Direct Blue 199 (I) and an ink containing 1.5 parts I were used for jet-printing resulting in clear images with improved optical d. after light irradiation

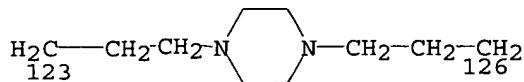
MSTR 3



G9 = 4-66 61-68

G14—G11—G14
4 61

G10 = alkenyl (opt. substd. by 1 or more G13)
 G11 = 123-4 126-61



G13 = CO2H
 G14 = 74

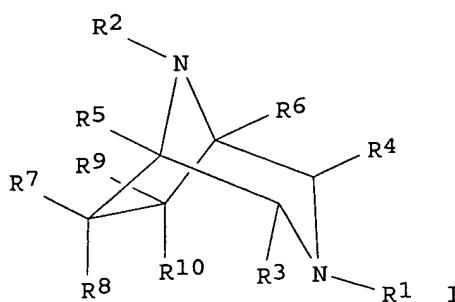
N—G10
74

Patent location: claim 9
 Note: or salts
 Note: additional ring formation also claimed

L14 ANSWER 11 OF 16 MARPAT COPYRIGHT 2006 ACS on STN
 ACCESSION NUMBER: 136:340701 MARPAT
 TITLE: Preparation of 3,8-diazabicyclo[3.2.1]octanes for treating cardiac arrhythmias
 INVENTOR(S): Bjoersne, Magnus; Hoffmann, Kurt-Juergen; Ponten, Fritiof; Strandlund, Gert; Svensson, Peder; Wilstermann, Michael
 PATENT ASSIGNEE(S): Astrazeneca AB, Swed.
 SOURCE: PCT Int. Appl., 135 pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

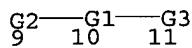
PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2002032902	A1	20020425	WO 2001-SE2294	20011018
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
AU 2001096173	A5	20020429	AU 2001-96173	20011018
EP 1328526	A1	20030723	EP 2001-977023	20011018
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR				
JP 2004511559	T2	20040415	JP 2002-536284	20011018
US 2004023971	A1	20040205	US 2003-399663	20030508
US 7012074	B2	20060314		
PRIORITY APPLN. INFO.:			SE 2000-3795	20001020
			WO 2001-SE2294	20011018

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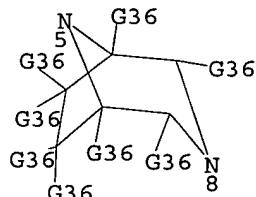


AB The title compds. [I; one of R1 and R2 = R1a and the other = ACR13R14BR15 (wherein R1a = alkyl optionally substituted and/or terminated by one or more groups selected from halo, CN, NO₂, etc.; R13 = H, halo, alkyl, etc.; R13R14 = O; or R14 = H, alkyl; R15 = (un)substituted aryl, heteroaryl; A = alkylene, etc.; B = a bond, alkylene, etc.); R3-R10 = H, alkyl], useful in the prophylaxis and in the treatment of arrhythmias, in particular atrial and ventricular arrhythmias, were prepared. Thus, reacting tert-Bu 3-(4-cyanophenoxy)-1-(3,8-diazabicyclo[3.2.1]oct-8-ylmethyl)propylcarbamate (preparation given) with Bu isocyanate in the presence of Et₃N in MeCN followed by treatment with HCl/EtOAc afforded I [R1 = CONHBu; R2 = CH₂CHNH₂CH₂CH₂O-p-C₆H₄CN; R3-R10 = H] in quant. yield. The exemplified compds. I showed pIC₅₀ values of at least 5.5 for K channel blockade.

MSTR 1

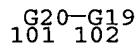


G₁ = 5-9 8-11



G₂ = carbon chain <containing 1-12 C>
(opt. subst. by 1 or more G46)

G₃ = 101



G₅ = carbon chain <containing 1-6 C> (opt. subst.)
G₁₁ = carbon chain <containing 1-6 C> (opt. subst.)
G₁₈ = carbon chain <containing 1-6 C> (opt. subst.)
G₂₀ = 125-10 126-102 / 127-10 129-102 /
103-10 105-102 / 121-10 124-102 / 141-10 143-102 /

148-10 151-102

$\begin{matrix} G_{11}-G_{21} \\ 125 \quad 126 \end{matrix}$ $\begin{matrix} G_{11}-G_{31}-G_{21} \\ 127 \quad 129 \end{matrix}$ $\begin{matrix} G_{11}-G_{21}-G_{24} \\ 103 \quad 104 \quad 105 \end{matrix}$ $\begin{matrix} G_{11}-G_{31}-G_{21}-G_{25} \\ 121 \quad 123 \quad 124 \end{matrix}$

$\begin{matrix} G_{11}-G_{21}-G_{32} \\ 141 \quad 142 \quad 143 \end{matrix}$ $\begin{matrix} G_{11}-G_{31}-G_{21}-G_{33} \\ 148 \quad 150 \quad 151 \end{matrix}$

G31 = 109

$\begin{matrix} N-G_{18} \\ 109 \end{matrix}$

G46 = 243

```

    graph LR
      N((N)) --- G5[G5]
      N --- G47[G47]
  
```

Patent location:

claim 1

Note: additional derivatization also claimed

Note: or pharmaceutically acceptable derivatives

Note: substitution is restricted

Note: also incorporates claims 36

REFERENCE COUNT: 7 THERE ARE 7 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L14 ANSWER 12 OF 16 MARPAT COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 134:193345 MARPAT

TITLE: Preparation of 4-amino-1-benzylpiperidines as antimalarials.

INVENTOR(S): Kim, Jin Mi; Ellman, Jonathan A.; Goldberg, Daniel

PATENT ASSIGNEE(S): Regents of the University of California, USA

SOURCE: PCT Int. Appl., 91 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2001014331	A2	20010301	WO 2000-US23338	20000823
WO 2001014331	A3	20010907		
W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM			
RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG			

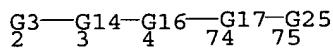
PRIORITY APPLN. INFO.:
GI

US 1999-150501P 19990824

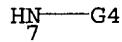
* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT *

AB R1R2R3 [R1 = R4X1(R5X2)NX6, Q1, Q2; R2 = (substituted) arylene, heteroarylene, alkylene; R3 = Q3, Q4; X1-X7 = bond, O, CO, CO2, CONH, CONR14, SO2; R4, R5, R8, R9, R11 = H, OH, alkoxy, (substituted) alkyl; R6, R7, R10, R12, R13 = CO, CO2, CONH, CONR15, SO2; R14-R17 = alkyl, aryl, heteroaryl, carboxylic acid ester or amide, amino, acylamino, alkoxy, OH, SH, phosphono, sulfonol, were prepared as inhibitors of protozoal proteases (no data). Thus, Cl(CH₂)₃Br and 4-amino-1-benzylpiperidine were stirred together for 16 h in MeCN to give 97% 1-benzyl-4-(3-chloropropylamino)piperidine. This was treated with di-tert-Bu dicarbonate in THF to give 94% 1-benzyl-4-(N-3-chloropropyl-N-tert-butoxycarbonylamino)piperidine. The latter was heated with NaN₃ and NaI in DMF at 75-80° to give 94% 1-benzyl-4-(N-3-azidopropyl-N-tert-butoxycarbonylamino)piperidine. Reduction with SnCl₂/PhSH/Et₃N in THF gave 93% 1-benzyl-4-(N-3-aminopropyl-N-tert-butoxycarbonylamino)piperidine. This was acylated with 4-benzyloxy-3,5-dimethoxybenzoic acid using PyBOP, HOAt, and DIPEA in DMF (88%) followed by deprotection with CF₃CO₂H in CH₂Cl₂ (94%) and reduction with LiAlH₄ in THF to give 64% title compound (I).

MSTR 1B

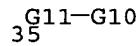


G3 = 7



G4 = carbon chain <containing 1-10 C,
0 or more double bonds, 0 or more triple bonds>
(opt. substd.)

G7 = 35



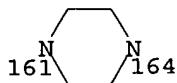
G10 = carbon chain <containing 1-10 C,
0 or more double bonds, 0 or more triple bonds>
(opt. substd.)

G11 = NH (opt. substd.)

G14 = carbon chain <containing 1-10 C,
0 or more double bonds, 0 or more triple bonds>
(opt. substd.)

G16 = bond

G17 = 161-4 164-75



G25 = 183

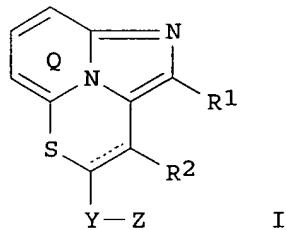
^{C(O)G7}
183

Patent location: claim 1
Note: substitution is restricted

L14 ANSWER 13 OF 16 MARPAT COPYRIGHT 2006 ACS on STN
 ACCESSION NUMBER: 130:311794 MARPAT
 TITLE: Preparation of fused imidazopyridine derivatives as hypolipemics and hypoglycemics
 INVENTOR(S): Takatani, Muneo; Sugiyama, Yasuo; Kawamoto, Tetsuji; Adachi, Koji
 PATENT ASSIGNEE(S): Takeda Chemical Industries, Ltd., Japan
 SOURCE: PCT Int. Appl., 153 pp.
 CODEN: PIIXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: Japanese
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 9920632	A1	19990429	WO 1998-JP4787	19981022
W: AL, AM, AU, AZ, BA, BB, BG, BR, BY, CA, CN, CU, CZ, EE, GD, GE, HR, HU, ID, IL, IS, JP, KG, KR, KZ, LC, LK, LR, LT, LV, MD, MG, MK, MN, MX, NO, NZ, PL, RO, RU, SG, SI, SK, SL, TJ, TM, TR, TT, UA, US, UZ, VN, YU, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
RW: GH, GM, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
AU 9896465	A1	19990510	AU 1998-96465	19981022
JP 11199586	A2	19990727	JP 1998-300748	19981022
PRIORITY APPLN. INFO.:			JP 1997-291023	19971023
			WO 1998-JP4787	19981022

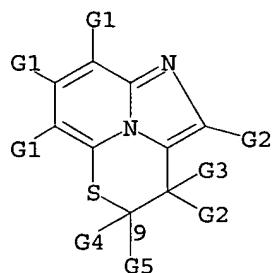
GI



AB The title compds. I [ring Q represents an optionally substituted pyridine ring; one of R1 and R2 represents hydrogen while the other represents

optionally substituted lower alkyl; Y represents a bond or optionally substituted divalent hydrocarbon group; Z represents a basic group optionally mediated by oxygen, nitrogen, CO, CS, SO₂N(R₃) (wherein R₃ represents hydrogen or optionally substituted hydrocarbyl) or S(O)_n (wherein n is 0, 1 or 2); and dotted line represents a single or double bond] are prepared I have the effects of increasing the low-d. lipoprotein receptor content, lowering the blood lipid level, and lowering the blood sugar level. The non-HDL cholesterol level in hamsters dosed with N-[1-(3-phenylpropan-1-yl)piperidin-4-yl]-2-methyl-5-thia-1,8b-diazacacenaphthylene-4-carboxamide dihydrochloride at 20 mg/kg/day for 4 days was 68.5% that of controls.

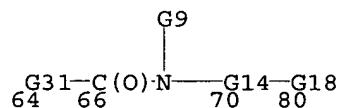
MSTR 1



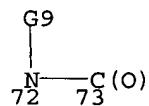
G5 = 19

₁₉^{G8}—₂₀^{G11}—₂₉^{G15}—₂₉^{G16}

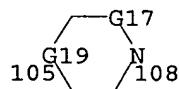
G8 = 64-9 80-20



G9 = alkenyl <containing 2-18 C>
 G11 = carbon chain (opt. substd.)
 G14 = carbon chain (opt. substd.)
 G15 = 72-20 73-29



G17 = (0-2) CH₂
 G18 = 105-70 108-20



G19 = N
 G31 = bond
 Derivative: or salts
 Patent location: claim 1
 Note: also incorporates claim 22
 Note: interruptions also claimed

REFERENCE COUNT: 16 THERE ARE 16 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L14 ANSWER 14 OF 16 MARPAT COPYRIGHT 2006 ACS on STN
 ACCESSION NUMBER: 128:266943 MARPAT
 TITLE: Piperazine-based cytofectins
 INVENTOR(S): Wheeler, Carl J.
 PATENT ASSIGNEE(S): Vical Inc., USA; Wheeler, Carl J.
 SOURCE: PCT Int. Appl., 44 pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 9814439	A1	19980409	WO 1997-US17155	19970924
W: CA, JP, US				
RW: AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE				
US 5861397	A	19990119	US 1996-726348	19961003
CA 2266781	AA	19980409	CA 1997-2266781	19970924
EP 929536	A1	19990721	EP 1997-943555	19970924
EP 929536	B1	20041222		
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, FI				
JP 2001511113	T2	20010807	JP 1998-516656	19970924
AT 285404	E	20050115	AT 1997-943555	19970924
US 6022874	A	20000208	US 1998-219758	19981223
PRIORITY APPLN. INFO.:			US 1996-726348	19961003
			WO 1997-US17155	19970924

AB The present invention relates to piperazine-based amphiphilic cationic lipids useful for gene therapy, transfection, and introducing immunogenic compds. for the purpose of vaccination. The disclosed compds. have lipophilic moieties linked to the ring nitrogens. In addition, at least one of the ring nitrogens is quaternized and linked to a hydrocarbon having at least one heteroatom. Numerous quaternized piperazines were prepared and employed both in vitro and in vivo for transfection of tumor, arterial and muscle cells. The effects of double quaternization of the piperazine moiety, of presence of heteroatoms (in the form of amino or hydroxyl groups) in the alkyl chains, and the alkyl chain length were studied.

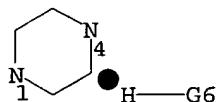
MSTR 1

G2—₇G1—G5—₉G1—G2

G1 = (1-6) CH₂
 G2 = 14

^{G3—G4}
14

G3 = NH
 G4 = alkenyl <containing 2-23 C, 1-6 double bonds>
 (opt. substd. by 1 or more G7)
 G5 = 1-7 4-9

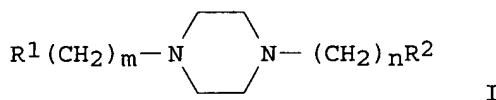


Patent location: claim 1
 Note: substitution is restricted
 Note: also incorporates claim 18

REFERENCE COUNT: 5 THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L14 ANSWER 15 OF 16 MARPAT COPYRIGHT 2006 ACS on STN
 ACCESSION NUMBER: 127:351188 MARPAT
 TITLE: Piperazine derivatives and drugs carriers containing them
 INVENTOR(S): Isozaki, Masafumi; Koiwai, Kazutomo; Uchiyama, Hideki
 PATENT ASSIGNEE(S): Terumo Corp., Japan
 SOURCE: Jpn. Kokai Tokkyo Koho, 7 pp.
 CODEN: JKXXAF
 DOCUMENT TYPE: Patent
 LANGUAGE: Japanese
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

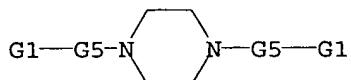
PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 09263582	A2	19971007	JP 1996-76331	19960329
PRIORITY APPLN. INFO.:			JP 1996-76331	19960329
GI				



AB The derivs. I ($\text{R}^{1-2} = \text{NR}^3\text{R}^4, \text{N}+\text{R}^5\text{R}^6\text{R}^7$; $\text{R}^3, \text{R}^5 = \text{C10-40 alkyl, alkenyl}; \text{R}^4, \text{R}^6-7 = \text{H, alkyl, alkenyl}$; $m, n = 1-10$) are claimed as components of drug carriers for diagnostic agents and/or therapeutic agents including DNA. The drug carriers may be in the forms of macromols., microaggregates, microparticles, microspheres, nanospheres, liposomes, or emulsions. The carriers are useful for effective transfer of nucleic acids, polynucleotide, gene, etc., to target tissues. A CHCl_3 solution containing 1,4-bis[3-(N-hexadecylamino)propyl]piperazine (preparation given), dilaurylphosphatidylcholine, and dioleoylphosphatidylethanolamine was evaporated in a flask and the flask was treated with an aqueous solution of plasmid

pcDNA/Amp bearing β -galactosidase gene under vigorous stirring to give liposomes. Transfection efficiency of the gene by the liposomes to Cos-1 cells was 17.8%, vs. 5.9% by Lipofectin.

MSTR 1



G1 = 11

$\frac{HN}{11}$ — G2

G2 = alkenyl <containing 1-40 C>

G5 = (1-10) CH₂

Patent location: claim 1

L14 ANSWER 16 OF 16 MARPAT COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 111:97286 MARPAT

TITLE: 4-[(Alkylamino)alkyl]piperazine derivatives as glutamic acid blockers

INVENTOR(S): Shinozaki, Atsuhiko; Sato, Masaru; Morifuji, Naoya; Hashimoto, Koichi; Kamishiro, Toshiro; Mazaki, Mitsuo

PATENT ASSIGNEE(S): Nippon Chemiphar Co., Ltd., Japan

SOURCE: Jpn. Kokai Tokkyo Koho, 9 pp.

CODEN: JKXXAF

DOCUMENT TYPE: Patent

LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

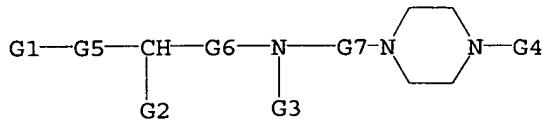
PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 01038080	A2	19890208	JP 1987-193202	19870801
PRIORITY APPLN. INFO.:			JP 1987-193202	19870801

GI For diagram(s), see printed CA Issue.

AB Alkylenediamine derivs. [I; R₁ = straight-chain or branched C₃-8 aliphatic hydrocarbyl, C₅-8 alicyclic hydrocarbyl, aryl, aryl-C₁-4 alkyl; R₂ = straight-chain or branched C₃-11 aliphatic hydrocarbyl, C₃-11 alkoxy, C₃-11 aliphatic hydrocarbyl containing an ester bond or an ether bond, aryloxy; R₃ = straight-chain, branched, or ester bond-containing C₃-11 aliphatic hydrocarbyl, aryl-ether bond-containing alkyl; when one of R₂ and R₃ is as described above, the other group is H or C_{≤2} alkyl; R₄ = H, straight-chain or branched C_{≤12} alkyl, C_{≤12} alkoxy, -acyl, or -alkoxycarbonyl, (un)substituted aryl, (un)substituted aryl-C₁-5 alkyl, C₁-3 hydroxyalkyl, cyano, R₁(CH₂)_mCHR₂(CH₂)_nNR₃(CH₂)_p; m, n = 0-3 integer; m + n <3; p = 2-13 integer] were prepared as CNS agents and insecticides. A solution of N-[4-methyl-1-(3-methylbutyl)pentyl]acrylamide 2.26 and 1-benzylpiperazine 2.12 g in MeOH was refluxed for 10 h to give 85% 3-(4-benzylpiperazinyl)-N-[4-methyl-1-(3-methylbutyl)pentyl]propanamide which was reduced by diborane in refluxing THF to give 71% 1-benzyl-4-[3-[4-methyl-1-(3-methylbutyl)pentylamino]propyl]piperazine. The latter in vitro reduced 87% glutamic acid-induced muscle-membrane electronic potential in the opener muscle of the dactyl in the first leg of crayfish (Combarus

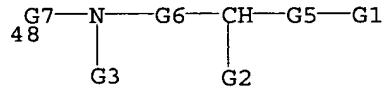
clarkii).

MSTR 1A



G3 = carbon chain <containing 3-11 C>
(opt. subst. by G8)

G4 = 48



G5 = bond

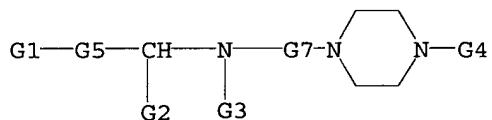
G7 = alkylene <containing 2-13 C, unbranched>

Derivative: or salts

Patent location: claims

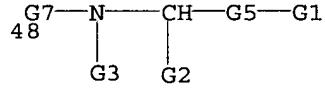
Note: substitution is restricted

MSTR 1B



G3 = carbon chain <containing 3-11 C>
(opt. subst. by G8)

G4 = 48



G5 = bond

G7 = alkylene <containing 2-13 C, unbranched>

Derivative: or salts

Patent location: claims

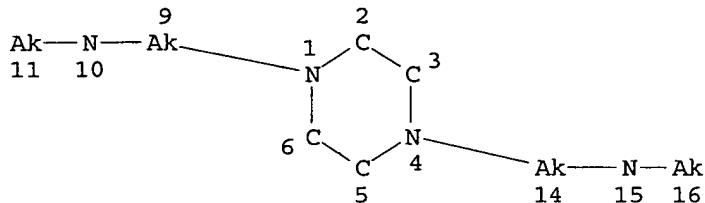
Note: substitution is restricted

FILE 'HOME' ENTERED AT 10:18:39 ON 31 MAR 2006

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L6 STR



NODE ATTRIBUTES:

CONNECT IS E2 RC AT 9
CONNECT IS E1 RC AT 11
CONNECT IS E2 RC AT 14
CONNECT IS E1 RC AT 16
DEFAULT MLEVEL IS ATOM
MLEVEL IS CLASS AT 9 11 14 16
GGCAT IS LIN LOC SAT AT 9
GGCAT IS UNS AT 11
GGCAT IS LIN LOC SAT AT 14
GGCAT IS UNS AT 16
DEFAULT ECLEVEL IS LIMITED
ECOUNT IS M6 C AT 11
ECOUNT IS M6 C AT 16

GRAPH ATTRIBUTES:

RING(S) ARE ISOLATED OR EMBEDDED
NUMBER OF NODES IS 12

STEREO ATTRIBUTES: NONE

L8 1 SEA FILE=REGISTRY SSS FUL L6

100.0% PROCESSED 706610 ITERATIONS
SEARCH TIME: 00.00.09

1 ANSWERS

(FILE 'HOME' ENTERED AT 10:01:52 ON 31 MAR 2006)

FILE 'CAPLUS' ENTERED AT 10:02:06 ON 31 MAR 2006

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E US2003-621760/AP,PRN 25
SET NOTICE 1000 SEARCH
L1 3 SEA ABB=ON US2003-621760/AP
SET NOTICE LOGIN SEARCH
SET LINE LOGIN
SET DETAIL LOGIN
D SCAN
SEL RN

FILE 'REGISTRY' ENTERED AT 10:02:56 ON 31 MAR 2006

L2 53 SEA ABB=ON (112-13-0/BI OR 112-16-3/BI OR 112-64-1/BI OR
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D SCAN

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FILE 'REGISTRY' ENTERED AT 10:06:35 ON 31 MAR 2006

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L6 STR L4
L7 0 SEA SSS SAM L6
L8 1 SEA SSS FUL L6
SAVE TEMP L8 POP760FULL/A
D SCAN
D LC

FILE 'REGISTRY' ENTERED AT 10:13:53 ON 31 MAR 2006
D STAT QUE L8

FILE 'CAPLUS, USPATFULL, TOXCENTER' ENTERED AT 10:13:53 ON 31 MAR 2006

L9 4 SEA ABB=ON L8
L10 2 DUP REM L9 (2 DUPLICATES REMOVED)
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ANSWER '2' FROM FILE USPATFULL
D IBIB ED ABS HITSTR 1-2

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L12 15 SEA SSS FUL L6
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FILE 'REGISTRY' ENTERED AT 10:17:03 ON 31 MAR 2006
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FILE 'CAPLUS, USPATFULL, TOXCENTER' ENTERED AT 10:17:03 ON 31 MAR 2006

L13 4 SEA ABB=ON L8

FILE 'MARPAT' ENTERED AT 10:17:20 ON 31 MAR 2006
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L14 16 DUP REM L13 L12 (3 DUPLICATES REMOVED)
ANSWER '1' FROM FILE CAPLUS
ANSWER '2' FROM FILE USPATFULL
ANSWERS '3-16' FROM FILE MARPAT
D IBIB ED ABS HITSTR 1-2

Popa 10/621760 search history

Page 3

D IBIB ED ABS QHIT 3-16

FILE 'HOME' ENTERED AT 10:18:39 ON 31 MAR 2006
D STAT QUE L8

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